

=> b hcap

FILE 'HCAPLUS' ENTERED AT 15:27:58 ON 21 AUG 2006

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FILE COVERS 1907 - 21 Aug 2006 VOL 145 ISS 9

FILE LAST UPDATED: 20 Aug 2006 (20060820/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d all hitstr l24 tot

L24 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2006 ACS on STN

AN 2004:289219 HCAPLUS

DN 140:273048

ED Entered STN: 08 Apr 2004

TI Procedure for the conversion of polysulfane in hydrogen sulfide and sulfur in gas flows resulting in hydrogen sulfide synthesis

IN Moeller, Alexander; Boeck, Wolfgang; Taugner, Wolfgang
; Heinzl, Harald; Rautenberg, Stephan

PA Degussa A.-G., Germany

SO Ger. Offen., 2 pp.

CODEN: GWXXBX

DT Patent

LA German

IC ICM B01D-0053/48

ICS B01D-0053/86

CC 49-2 (Industrial Inorganic Chemicals)

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
PI	DE--10245164	A1	20040408	2002DE-1045164	20020926 <--	
	WO2004028963	A1	20040408	2003WO-EP09432	20030826 <--	
	W:			AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW		
	RW:			GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG		
	AU2003255483	A1	20040419	2003AU-0255483	20030826 <--	
	EP---1542925	A1	20050622	2003EP-0798130	20030826 <--	
	R:			AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK		
	BR2003014663	A	20050802	2003BR-0014663	20030826 <--	
	CN---1684905	A	20051019	2003CN-0823024	20030826 <--	
	JP2006500309	T2	20060105	2004JP-0538838	20030826 <--	
	US2005265913	A1	20051201	2005US-0529148	20050324 <--	

PRAI 2002DE-1045164 A 20020926 <--
 2003WO-EP09432 W 20030826

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
DE 10245164	ICM	B01D-0053/48
	ICS	B01D-0053/86
	IPCI	B01D0053-48 [ICM,7]; B01D0053-86 [ICS,7]
	IPCR	C01B0017-00 [I,C*]; C01B0017-16 [I,A]
	ECLA	C01B017/16M; C01B017/16P
WO2004028963	IPCI	C01B0017-16 [ICM,7]; C01B0017-00 [ICM,7,C*]
	IPCR	C01B0017-00 [I,C*]; C01B0017-16 [I,A]
	ECLA	C01B017/16M; C01B017/16P
AU2003255483	IPCI	C01B0017-16 [ICM,7]; C01B0017-00 [ICM,7,C*]
	IPCR	C01B0017-00 [I,C*]; C01B0017-16 [I,A]
EP---1542925	IPCI	C01B0017-16 [ICM,7]; C01B0017-00 [ICM,7,C*]
	IPCR	C01B0017-00 [I,C*]; C01B0017-16 [I,A]
	ECLA	C01B017/16M; C01B017/16P
BR2003014663	IPCI	C01B0017-16 [ICM,7]; C01B0017-00 [ICM,7,C*]
	IPCR	C01B0017-00 [I,C*]; C01B0017-16 [I,A]
	ECLA	C01B017/16M; C01B017/16P
CN---1684905	IPCI	C01B0017-16 [ICM,7]; C01B0017-00 [ICM,7,C*]
	IPCR	C01B0017-00 [I,C*]; C01B0017-16 [I,A]
JP2006500309	IPCI	C01B0017-16 [I,A]; C01B0017-00 [I,C*]
US2005265913	IPCI	C01B0017-20 [ICM,7]; C01B0017-00 [ICM,7,C*]
	NCL	423/242.200
	ECLA	C01B017/16M; C01B017/16P
AB	Polysulfane (H ₂ S _x) resulting in hydrogen sulfide synthesis are catalytically converted by contacting with e.g. activated carbon, Al ₂ O ₃ , SiO ₂ , or zeolithes to give H ₂ S and S.	
ST	polysulfane catalytic conversion; hydrogen sulfide manuf; sulfur manuf	
IT	Zeolites (synthetic), processes	
	RL: CAT (Catalyst use); CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)	
	(for conversion of polysulfane in hydrogen sulfide and sulfur in gas flows resulting in hydrogen sulfide synthesis)	
IT	7440-44-0, Carbon, processes	
	RL: CAT (Catalyst use); CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)	
	(activated; for conversion of polysulfane in hydrogen sulfide and sulfur in gas flows resulting in hydrogen sulfide synthesis)	
IT	1344-28-1, Alumina, processes 7631-86-9, Silica, processes	
	RL: CAT (Catalyst use); CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)	
	(for conversion of polysulfane in hydrogen sulfide and sulfur in gas flows resulting in hydrogen sulfide synthesis)	
IT	7704-34-9P, Sulfur, preparation 7783-06-4P, Hydrogen sulfide, preparation	
	RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)	
	(procedure for conversion of polysulfane in hydrogen sulfide and sulfur in gas flows resulting in hydrogen sulfide synthesis)	
IT	37331-50-3, Sulfane	
	RL: RCT (Reactant); RACT (Reactant or reagent)	
	(procedure for conversion of polysulfane in hydrogen sulfide and sulfur in gas flows resulting in hydrogen sulfide synthesis)	
IT	7783-06-4P, Hydrogen sulfide, preparation	
	RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)	
	(procedure for conversion of polysulfane in hydrogen sulfide and sulfur in gas flows resulting in hydrogen sulfide synthesis)	
RN	7783-06-4 HCAPLUS	
CN	Hydrogen sulfide (H ₂ S) (8CI, 9CI) (CA INDEX NAME)	

H₂S

IT 37331-50-3, Sulfane
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (procedure for conversion of polysulfane in hydrogen sulfide and sulfur
 in gas flows resulting in hydrogen sulfide synthesis)
 RN 37331-50-3 HCAPLUS
 CN Sulfane (9CI) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

L24 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2006 ACS on STN
 AN 1986:409243 HCAPLUS
 DN 105:9243
 ED Entered STN: 13 Jul 1986
 TI Removing sulfane from biogas
 IN Buryan, Petr; Zacher, Jan; Palaty, Jiri; Jonas, Jaroslav
 PA Czech.
 SO Czech., 2 pp.
 CODEN: CZXXA9
 DT Patent
 LA Czech
 IC C10K-0001/14
 CC 52-1 (Electrochemical, Radiational, and Thermal Energy Technology)
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI CS-----221457	B	19830429	1981CS-0001999	19810318
PRAI 1981CS-0001999		19810318		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
CS 221457	IC	C10K-0001/14
	IPCI	C10K0001-14; C10K0001-00 [C*]
	IPCR	C10K0001-00 [I,C*]; C10K0001-14 [I,A]

AB The content of sulfanes in biogas is decreased to 1% after scrubbing with aqueous FeL- (H₄L = EDTA) at 20-50°. The reduced form of the agent FeL₂- is reactivated by air oxidation

ST hydrogen sulfide removal biogas; iron EDTA hydrogen sulfide removal

IT Fuel gas manufacturing
 (biogas, hydrogen sulfide removal in, iron EDTA salt for)

IT 74-82-8P, preparation
 RL: PREP (Preparation)
 (manufacture of gas containing, hydrogen sulfide removal in, iron EDTA salt for)

IT 15275-07-7
 RL: USES (Uses)
 (removal of hydrogen sulfide, from biogas)

IT 7783-06-4, uses and miscellaneous
 RL: REM (Removal or disposal); PROC (Process)
 (removal of, from biogas, iron EDTA salt in)

L24 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2006 ACS on STN
 AN 1969:456115 HCAPLUS
 DN 71:56115
 ED Entered STN: 12 May 1984
 TI Sulfanes
 AU Burton, K. W. C.; Machmer, Paul
 CS Univ. Cologne, Cologne, Fed. Rep. Ger.
 SO Inorg. Sulphur Chem. (1968), 335-66. Editor(s): Nickless, G. Publisher: Elsevier Publ. Co., Amsterdam, Neth.
 CODEN: 21AEAH
 DT Conference; General Review
 LA English

CC 78 (Inorganic Chemicals and Reactions)
 AB Sulfanes are reviewed in terms of H₂S, preparation and phys. properties of sulfanes, thermal chemistry of polysulfanes, Raman and uv spectra of sulfanes, mol. distribution function of the sulfane-halosulfane condensation reaction, chemistry of sulfanes including reactions with chloral and Cl₃CSCl, and asym. derivs. of the type RSnH and RSmCl, thioalkanes, acidity of the sulfanes, and anal. determination of sulfanes and sulfane-S mixts. 150 references.
 ST sulfanes review; review sulfanes; anal
 IT sulfanes review; spectra sulfanes review
 IT 50864-71-6P, Hydrogen sulfide (H₂Sx)
 RL: SPN (Synthetic preparation); PREP (Preparation)

=> d his

(FILE 'HOME' ENTERED AT 15:06:04 ON 21 AUG 2006)

FILE 'HCAPLUS' ENTERED AT 15:06:51 ON 21 AUG 2006

L1 1 US2005265913/PN OR (US2005-529148 OR DE2002-10245164)/AP,PRN
 E MOLLER A/AU
 L2 81 E3-15
 L3 3 E24
 E MOELLER A/AU
 L4 121 E3-11,E24-25
 E BOCK W/AU
 L5 181 E3-7,E30-31
 E TAIGNER W/AU
 E TAUGNER W/AU
 L6 4 E4
 E HEINZEL H/AU
 L7 13 E3,E5
 E RAUTENBERG S/AU
 L8 7 E4-5

FILE 'REGISTRY' ENTERED AT 15:12:36 ON 21 AUG 2006

FILE 'HCAPLUS' ENTERED AT 15:12:36 ON 21 AUG 2006

L9 TRA L1 1- RN : 6 TERMS

FILE 'REGISTRY' ENTERED AT 15:12:36 ON 21 AUG 2006

L10 6 SEA L9
 L11 136 SULFANE
 L12 1230 H₂S

FILE 'HCAPLUS' ENTERED AT 15:16:58 ON 21 AUG 2006

L13 52463 L12
 L14 108157 ?HYDROGEN (1A) (SULFID? OR SULPHID?) OR (HYDROSULFUR? OR HYDROSU
 E HYDROGEN SULFIDE/CT
 E E3+ALL
 L15 50816 E4
 L16 7703 L13-15 (L) PREP+NT/RL
 L17 5 L16 AND L1-8
 L18 9019 L11
 L19 511 SULFANE OR SULPHANE
 E POLYSUFANE/CT
 E POLYSULFANE/CT
 L20 1 L17 AND L18-19
 L21 13 L16 AND L19
 L22 12 L21 NOT L20
 SEL AN 7 11
 L23 2 E1-4 AND L22
 L24 3 L20,L23
 L25 133 POLYSULFAN? OR POLYSULPHAN?
 L26 5 L25 AND L16

L27	1 L26 AND L1-8
L28	4 L26 NOT L27
L29	2 L28 NOT L22

=> b wpix

FILE 'WPIX' ENTERED AT 09:37:56 ON 22 AUG 2006
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FILE LAST UPDATED: 17 AUG 2006 <20060817/UP>
MOST RECENT DERWENT UPDATE: 200653 <200653/DW>
DERWENT WORLD PATENTS INDEX SUBSCRIBER FILE, COVERS 1963 TO DATE

>>> FOR A COPY OF THE DERWENT WORLD PATENTS INDEX STN USER GUIDE,
PLEASE VISIT:
http://www.stn-international.de/training_center/patents/stn_guide.pdf <

>>> FOR DETAILS OF THE PATENTS COVERED IN CURRENT UPDATES, SEE
<http://scientific.thomson.com/support/patents/coverage/latestupdates/>

>>> PLEASE BE AWARE OF THE NEW IPC REFORM IN 2006, SEE
http://www.stn-international.de/stdatabases/details/ipc_reform.html and
<http://scientific.thomson.com/media/scpdf/ipcrdwpf.pdf> <<<

>>> FOR FURTHER DETAILS ON THE FORTHCOMING DERWENT WORLD PATENTS
INDEX ENHANCEMENTS PLEASE VISIT:
http://www.stn-international.de/stdatabases/details/dwpi_r.html <<<
'BIX' IS DEFAULT SEARCH FIELD FOR 'WPIX' FILE

=> d all abex tech l30 tot

L30 ANSWER 1 OF 4 WPIX COPYRIGHT 2006 THE THOMSON CORP on STN

AN 2004-317818 [30] WPIX

DNC C2004-120857

TI Purification of crude gas stream from hydrogen sulfide
synthesis comprises catalytic conversion of polysulfanes into
hydrogen sulfide and sulfur.

DC E36 J01 J04

IN BOCK, W; HEINZEL, H; MOLLER, A;
RAUTENBERG, S; TAUGNER, W; BOECK, W; MOELLER, A

PA (DEGS) DEGUSSA AG; (BOCK-I) BOCK W; (HEIN-I) HEINZEL H; (MOLL-I) MOLLER A;
(RAUT-I) RAUTENBERG S; (TAUG-I) TAUGNER W

CYC 106

PI DE----10245164 A1 20040408 (200430)* 2 B01D-053-48

WO--2004028963 A1 20040408 (200430) EN C01B-017-16

RW: AT BE BG CH CY CZ DE DK EA EE ES FI FR GB GH GM GR HU IE IT KE LS

LU MC MW MZ NL OA PT RO SD SE SI SK SL SZ TR TZ UG ZM ZW

W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK

DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR

KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NI NO NZ OM PG PH

PL PT RO RU SC SD SE SG SK SL SY TJ TM TN TR TT TZ UA UG US UZ VC

VN YU ZA ZM ZW

AU--2003255483 A1 20040419 (200462) C01B-017-16

EP-----1542925 A1 20050622 (200541) EN C01B-017-16

R: AL AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HU IE IT LI LT LU LV

MC MK NL PT RO SE SI SK TR

BR---200314663 A 20050802 (200553) C01B-017-16

US--2005265913 A1 20051201 (200579) C01B-017-20 <--

JP--2006500309 W 20060105 (200603) 8 C01B-017-00

CN-----1684905 A 20051019 (200612) C01B-017-16

ADT DE----10245164 A1 2002DE-1045164 20020926; WO--2004028963 A1

2003WO-EP09432 20030826; AU--2003255483 A1 2003AU-0255483 20030826;

EP-----1542925 A1 2003EP-0798130 20030826, 2003WO-EP09432 20030826;

BR---200314663 A 2003BR-0014663 20030826, 2003WO-EP09432 20030826;

US--2005265913 A1 2003WO-EP09432 20030826, 2005US-0529148 20050324

; JP--2006500309 W 2003WO-EP09432 20030826, 2004JP-0538838 20030826;

CN-----1684905 A 2003CN-0823024 20030826

FDT AU--2003255483 A1 Based on WO--2004028963; EP-----1542925 A1 Based on

WO--2004028963; BR---200314663 A Based on WO--2004028963; JP--2006500309 W

Based on WO--2004028963

PRAI 2002DE-1045164 20020926

IC ICM B01D-053-48; C01B-017-00; C01B-017-16; C01B-017-20
ICS B01D-053-86

AB DE 10245164 A UPAB: 20040511

NOVELTY - Process for converting polysulfanes into hydrogen sulfide (H₂S) and sulfur (S).comprises catalytic conversion of polysulfanes (H₂S_x) in crude gas streams containing H₂S, obtained in H₂S synthesis.

USE - The process is used for purifying crude gas streams obtained in hydrogen sulfide synthesis.

ADVANTAGE - Crude gas obtained in synthesis of hydrogen sulfide (H₂S) from hydrogen and sulfur (S) usually contains polysulfanes (H₂S_x), in amounts over 400 vpm, as by-products. Compressing the gas tends to cause uncontrolled decomposition into H₂S and S and hence undesirable deposition of S in the entire compression zone, including the peripheral pipes and valves. The present process results in controlled conversion of polysulfanes and prevents deposition of S in the pipework.

Dwg.0/0

FS CPI

FA AB; DCN

MC CPI: E10-B03; E11-Q02; E31-D04; E31-F02; E31-N04C; E31-N04D; E31-P02B; E31-P03; E32-A02; E33-A03; E33-A04; J01-E03F; J04-E01; N01-A; N01-B; N01-C02; N01-D02; N02; N03; N04-A; N05-D; N06-A; N06-B; N07-L02B

ABEX UPTX: 20040511

EXAMPLE - Crude hydrogen sulfide (H₂S) gas (up to 5000 Nm³/hour), generated from hydrogen (H₂) and sulfur (S), was passed at 1.05-1.5 bar absolute into a jet scrubber system for converting the polysulfanes into H₂S and S. The scrubbing liquid was an aqueous or methanolic 0.5-10% solution of potassium and/or sodium hydroxide/hydrogen sulfide (KOH/KHS, NaOH/NaHS). The resultant S remained in solution as the corresponding polysulfide. Any precipitated solid S could be removed by filtration. Fresh solution was used to replace losses by evaporation and the fraction of circulating solution discharged, according to the S content. More sulfane was decomposed in a countercurrent scrubber and droplets were separated in a demister. If necessary, residual sulfanes were decomposed in an adsorber bed (activated charcoal, zeolite etc.) and the S formed was deposited. Online UV analysis was used to determine the sulfane concentration in the crude and purified gas. The process prevented undesired S deposits in the plant.

TECH UPTX: 20040511

TECHNOLOGY FOCUS - CHEMICAL ENGINEERING - Preferred Process: The crude gas containing H₂S is contacted with (a) a suitable catalytic solid; (b) a basic aqueous or alcoholic solution containing catalytically active compounds; or (c) a gas containing catalytically active compounds. Conversion may be carried out in several stages.

TECHNOLOGY FOCUS - INORGANIC CHEMISTRY - Catalysts: (disclosed) Suitable catalysts are (a) activated charcoal, alumina, silica in various forms, including naturally-occurring minerals, zeolites, glasses, (mixed) oxides, alkali(ne earth) and other basic (hydr)oxide(s); (b) solutions of ammonia, amines, aminoalcohols and alkali(ne earth) or other basic (hydr)oxides or (hydrogen) sulfides; (c) gaseous ammonia, amines or aminoalcohols.

L30 ANSWER 2 OF 4 WPIX COPYRIGHT 2006 THE THOMSON CORP on STN

AN 2004-259406 [25] WPIX

DNC C2004-101338

TI Purification of hydrogen sulphide synthesised from hydrogen and liquid sulfur by passage through a porous filter compound.

DC E36 J01

IN LE BEC, R; LE BEC, R O M

PA (AQOR) ATOFINA; (ARKE-N) ARKEMA; (ARKE-N) ARKEMA INC; (AQOR) ARKEMA; (AQOR) ATOFINA SA

CYC 106

PI FR-----2844208 A1 20040312 (200425)* 9 B01D-046-00
 WO--2004022482 A2 20040318 (200425) FR C01B-017-16
 RW: AT BE BG CH CY CZ DE DK EA EE ES FI FR GB GH GM GR HU IE IT KE LS
 LU MC MW MZ NL OA PT RO SD SE SI SK SL SZ TR TZ UG ZM ZW
 W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK
 DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR
 KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NI NO NZ OM PG PH
 PL PT RO RU SC SD SE SG SK SL SY TJ TM TN TR TT TZ UA UG US UZ VC
 VN YU ZA ZM ZW

AU--2003278253 A1 20040329 (200459) C01B-017-16

EP-----1542924 A2 20050622 (200541) FR C01B-017-16

R: AL AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HU IE IT LI LT LU LV
 MC MK NL PT RO SE SI SK TR

EP-----1542924 B1 20060201 (200612) FR C01B-017-00

R: AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HU IE IT LI LU MC NL PT
 RO SE SI SK TR

DE----60303453 E 20060413 (200629) C01B-017-00

ES-----2257696 T3 20060801 (200652) C01B-017-16

ADT FR-----2844208 A1 2002FR-0011156 20020906; WO--2004022482 A2
 2003WO-FR002638 20030903; AU--2003278253 A1 2003AU-0278253 20030903;
 EP-----1542924 A2 2003EP-0769563 20030903, 2003WO-FR02638 20030903;
 EP-----1542924 B1 2003EP-0769563 20030903, 2003WO-FR02638 20030903;
 DE----60303453 E 2003DE-0603453 20030903, 2003EP-0769563 20030903,
 2003WO-FR02638 20030903; ES-----2257696 T3 2003EP-0769563 20030903
 FDT AU--2003278253 A1 Based on WO--2004022482; EP-----1542924 A2 Based on
 WO--2004022482; EP-----1542924 B1 Based on WO--2004022482; DE----60303453
 E Based on EP-----1542924, Based on WO--2004022482; ES-----2257696 T3
 Based on EP-----1542924

PRAI 2002FR-0011156 20020906

IC ICM B01D-046-00; C01B-017-00; C01B-017-16

ICS B01J-020-08; B01J-020-10; B01J-020-20

AB FR 2844208 A UPAB: 20040418

NOVELTY - Purification of a gas comprising mainly hydrogen sulfide obtained by reaction of hydrogen with liquid sulfur in an industrial unit, by passing the gas through a filter containing porous particles of active carbon, alumina or silica.

USE - The process of the invention resolves the problem of formation and decomposition of sulfanes, H₂S_x, where x is 2 or more.

ADVANTAGE - The advantage of using porous materials is that they become saturated with sulfur and/or sulfur compounds in the interior of their pores, thus avoiding blockage in the spaces between the particles. Active carbon is capable of retaining up to 70% of its initial weight in sulfurised compounds. After use, the used carbon can be incinerated and converted entirely to CO, SO₂ and H₂O.

Dwg.0/0

FS CPI

FA AB; DCN

MC CPI: E11-Q01; E31-F02; E31-N04C; E31-P03; E34-C02; J01-G03

ABEX UPTX: 20040418

EXAMPLE - A filter containing active carbon was submitted to a current of gas, of purity 99.7% in H₂S, for a determined time. The filter was then isolated from the gas current and purged with N₂ at 20 - 100 degrees C to eliminate H₂S. The direction of traverse of the filter defined an entry and exit, and samples of active carbon were taken at regularly spaced intervals between entry and exit for analysis. Total sulfur was determined by microanalysis; the sample was submitted to total combustion in the presence of O₂, the S compounds being converted to SO₂ then H₂SO₄ by oxidation with H₂O₂ and estimation by coulometry. A synthesis gas, at 4 bars pressure, containing mainly H₂S from a S/H₂ reaction was passed through a condenser cooling to 30 degrees C at 0.5 tonne/hour (75 cm³/hour). The gas was then passed through a cylindrical filter containing 8 kg of active carbon, ACTICARBONE AC35. The active carbon was present as cylinders of 4 mm diameter and surface area at least 1000 m².g. After 8 hours of filtration, the filter was removed and analysed; samples were taken at spacings of 10 cm along the length of the

filter from 0(entry), 10, 20, 30, 40 and 50(exit).The results, in this order, as total S in g/100 g of initial active carbon were :- 0(34), 10(24), 20(1), 30(1), 40(1) and 50(1).

TECH UPTX: 20040418

TECHNOLOGY FOCUS - INORGANIC CHEMISTRY - Preferred Process Details : The industrial process is carried out at 400 - 450 degrees C in a reactor over mounted with a reflux column ; the gas leaving the head of the column is cooled in one or more condensers where sulfur is recovered. The gas leaving the condensers, at 30 degrees C, may contain impurities leading to a post reaction in which sulfanes, H₂Sx, as well as sulfur, can cause blockages and product breakdowns. The porous filter compound is preferably active carbon ; the filter may also contain a material for selective adsorption of water such as a molecular sieve of type 3 Angstrom.

L30 ANSWER 3 OF 4 WPIX COPYRIGHT 2006 THE THOMSON CORP on STN

AN 1990-369027 [50] WPIX

DNC C1990-160546

TI Water treatment agent - comprises sulphide- and sulphate-ions and sulphur, etc., for heavy- and transition-metal removal.

DC D15 E37 M25

IN STRATEN, G

PA (STRA-I) STRATEN G

CYC 18

PI DE-----3917412 C 19901213 (199050)*

WO-----9200917 A 19920123 (199207)#

RW: AT BE CH DE DK ES FR GB IT LU NL SE

W: CA DK FI NO SU US

EP-----537143 A1 19930421 (199316)# GE 46 C02F-001-52

R: AT BE CH DE DK ES FR GB IT LI LU NL SE

US-----5451327 A 19950919 (199543)# 11 C02F-001-62

EP-----537143 B1 19970917 (199742)# GE 13 C02F-001-52

R: AT BE CH DE DK ES FR GB IT LI LU NL SE

DE----59010762 G 19971023 (199748)# C02F-001-52

ADT DE-----3917412 C 1989DE-3917412 19890529; EP-----537143 A1 1990EP-0910100

19900704, 1990WO-DE00499 19900704; US-----5451327 A 1990WO-DE00499

19900704, 1993US-0972479 19930318; EP-----537143 B1 1990EP-0910100

19900704, 1990WO-DE00499 19900704; DE----59010762 G 1990DE-0510762

19900704, 1990EP-0910100 19900704, 1990WO-DE00499 19900704

FDT EP-----537143 A1 Based on WO-----9200917; US-----5451327 A Based on

WO-----9200917; EP-----537143 B1 Based on WO-----9200917; DE----59010762

G Based on EP-----537143, Based on WO-----9200917

PRAI 1989DE-3917412 19890529; 1990EP-0910100 19900704;

1993US-0972479 19930318; 1990DE-0510762 19900704

REP 1.Jnl.Ref; BE----447850; DD----154008; EP----349671; US---1934626

IC ICM C02F-001-52; C02F-001-62

ICS C01B-017-22; C02F-001-58

AB DE 3917412 C UPAB: 19930928

Water treatment agent (I) comprises (A) 17-21% S22- ions; (B) 4-8% S32- ions; (C) 15-21% S42- ions; (D) 3-7% S52- ions; (E) 12-18% S62- ions; (F) 10-14% S82- ions; (G) 11-16% S2O32- ions; (H) 6-10% S4O32- ions; (I) 1-5% S4O62- ions, and (J) 0-3% S6 and ring sulphur.

USE/ADVANTAGE - For the removal of heavy or transition metals from water. Unlike conventional metal hydroxide treatment solns., (I) can be used over a wide pH range and produces heavy metal ppts. that are easily removed. (I) can especially be used with spent corrosion inhibitors or waste streams from metal alloy mfr.

0/3

FS CPI

FA AB; DCN

MC CPI: D04-A01B; D04-B05; E31-F04; E31-F05; M25-E01

L30 ANSWER 4 OF 4 WPIX COPYRIGHT 2006 THE THOMSON CORP on STN

AN 1989-301845 [42] WPIX

DNC C1989-133437

TI Sealing of gas-separation membranes, especially hollow fibres - by surface treatment

with pref. di-or tri-functional monomers inter-reacting to form polymers.

DC A28 A88 E36 J01 P42
 IN EKINER, O M; HAYES, R A; MANOS, P; EKINER, O; EKINER, C M
 PA (AIRL) AIR LIQUIDE SA; (DUPO) DU PONT DE NEMOURS & CO E I; (CAAL) AIR
 LIQUIDE CANADA LTEE
 CYC 23
 PI EP-----336999 A 19891018 (198942)* EN 19
 R: AT BE CH DE ES FR GB GR IT LI LU NL SE
 JP-----01262922 A 19891019 (198948)
 NO-----8802144 A 19891106 (198950)
 PT-----87511 A 19891110 (198950)
 DK-----8802692 A 19891014 (198951)
 BR-----8802375 A 19891205 (199003)
 ZA-----8803525 A 19900131 (199009)
 AU-----8816330 A 19900412 (199023)
 CN-----1036908 A 19891108 (199033)
 US-----5091216 A 19920225 (199211) 11
 JP-----93077446 B 19931026 (199345) 15 B01D-071-82
 CA-----1329898 C 19940531 (199427) B01D-069-00
 EP-----336999 B1 19950906 (199540) EN 21 B01D-071-06
 R: AT BE CH DE ES FR GB GR IT LI LU NL SE
 DE-----3854426 G 19951012 (199546) B01D-071-06
 ADT EP-----336999 A 1988EP-0107943 19880518; JP-----01262922 A 1988JP-0119439
 19880518; ZA-----8803525 A 1988ZA-0003525 19880518; US-----5091216 A
 1990US-0622269 19901205; JP-----93077446 B 1988JP-0119439 19880518;
 CA-----1329898 C 1988CA-0567000 19880517; EP-----336999 B1 1988EP-0107943
 19880518; DE-----3854426 G 1988DE-3854426 19880518, 1988EP-0107943
 19880518
 FDT JP-----93077446 B Based on JP-----01262922; DE-----3854426 G Based on
 EP-----336999
 PRAI 1988US-0175499 19880413
 REP 2.Jnl.Ref; A3...9135; FR---2391752; JP--59059220; JP--59059222; No-SR.Pub;
 WO---8810140; JP---5959220; JP---5959222
 IC B01D-013-04; B01D-053-22; B05D-005-00; C08J-005-22; C08J-007-16
 ICM B01D-069-00; B01D-071-06; B01D-071-82
 ICS B01D-013-04; B01D-053-22; B05D-005-00; C08J-005-22; C08J-007-16;
 D06M-013-152; D06M-013-165; D06M-013-192; D06M-013-332; D06M-013-395
 AB EP 336999 A UPAB: 19930923
 Sealing of a gas-separation membrane is effected by applying to the surface at
 least two monomers which can react to form a polymer and thus improve the
 membrane selectivity. The monomer combinations are pref. of di- or
 tri-functional monomers and pref. comprise (A) an acylchloride or an
 isocyanate or a glycidyl ether and (B) an amine.
 USE/ADVANTAGE - The method is pref. used with aromatic polyamide or
 aromatic polysulphane membranes which are in hollow fibre form
 (claimed) e.g. of the type described in US4230463. The polymer formed by
 (A) and (B) serves to seal defects or imperfections arising during
 membrane formation or during subsequent membrane handling. Among the
 membranes thus treated are those used in H2 recovery from refinery or
 ammonia plant; separation of CO or H2S from hydrocarbons; or enrichment of O2
 and N2 from air.
 O/O
 FS CPI GMPI
 FA AB; DCN
 MC CPI: A11-B05C; A12-S05A; A12-W11A; E10-J02D; E11-Q01; E31-A02; E31-D01;
 E31-F02; E31-H04; E31-N05B; J01-C03; J01-E03E

=> => d his

(FILE 'HOME' ENTERED AT 09:17:07 ON 22 AUG 2006)

FILE 'REGISTRY' ENTERED AT 09:17:40 ON 22 AUG 2006

L1 1 HYDROGEN SULFIDE/CN

FILE 'WPIX' ENTERED AT 09:18:14 ON 22 AUG 2006

L2 1 US2005265913/PN OR (US2005-529148 OR DE2002-10245164)/AP, PRN
E MOLLER A/AU
L3 86 E3-7
E MOELLER A/AU
L4 95 E3-6
E BOCK W/AU
L5 92 E3-4
E TAUGNER W/AU
L6 4 E3
E HEINZEL H/AU
L7 11 E3
E RAUTENBERG S/AU
L8 8 E3
E HYDROGEN SULFIDE/CN
L9 2 E3-4
L10 7244 (129436-0-0-0 OR 357-0-0-0)/DCRE OR (R01785 OR RA01M1)/DCN OR 1
L11 1526 L9/DCR
L12 18915 (DIHYDROGEN OR HYDROGEN) (1A) (?SULPHID? OR ?SULFID?) OR STINK DA
L13 20295 L10-12
L14 158 POLYSULFANE? OR POLYSULPHANE? OR SULFANE? OR SULPHANE?
L15 12 L13 AND L14
L16 1 L15 AND L2-8
L17 11 L15 NOT L16
L18 1688 E31-F02/MC
L19 3 L18 AND L14
L20 1 L19 AND L2-8
L21 3 L16, L19
L22 10 H2SX
L23 2 L22 AND L14
L24 1 L23 AND L2-8
L25 1 L23 NOT L24
L26 2 L19 NOT L20
L27 12 L17, L24, L26
L28 3 L21, L24
SEL AN 2 3 6 L27
L29 3 E1-3 AND L27
L30 4 L28-29

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